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AMENDMENTS TO THE CLAIMS

1. (currently amended) A method for the synthesis of severely sterically hindered secondary aminoether alcohols of the formula

wherein R¹ and R² are each selected from the group consisting of alkyl, hydroxylalkyl radicals having 1 to 4 carbon atoms or in combination with the carbon atom to which they are attached they form a cycloalkyl group having 3 to 8 carbon atoms, and R³ is selected from the group consisting of hydrogen, alkyl, hydroxyalkyl radicals having 1 to 4 carbon atoms, and mixtures thereof, and R⁴, R⁵, R⁶, R⁷, R⁸, R⁹, R¹⁰ and R¹¹ are the same or different and are selected from the group consisting of hydrogen, alkyl and hydroxyalkyl radicals having 1 to 4 carbons provided that at least one of R⁴ or R⁵ bonded to the carbon atom directly bonded to the nitrogen atom is an alkyl or hydroxyalkyl radical when R³ is hydrogen, the process involving reacting an organic carboxylic acid or salt of a carboxylic acid of the formula

$$R^{12} - C - OY$$

wherein R^{12} is selected from the group consisting of alkyl radicals having 1 to 4 carbon atoms, aryl radicals bearing hydrogen or one or more C_1 - C_{10} alkyl groups substituted thereon, and mixtures thereof, and Y is selected from the group consisting of hydrogen, alkali metal, ammonium, and mixtures thereof, with a sulfonyl halide, a sulfuryl halide, a mixed sulfuryl ester halide, or a mixed sulfuryl amide halide of the formula

$$R^{14}SO_2X$$
 , SO_2X_2 , $R^{14}OSO_2X$, or $R_2^{14,14}NSO_2X$

wherein X is selected from the group consisting of F, Cl, Br, I, and mixtures thereof, and R¹⁴ and R¹⁴ are the same or different and each is selected from the group consisting of alkyl radicals having 1 to 4 carbon atoms, haloalkyl radicals of the formula

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 $C_nH_{(2n+1)-w}Z_w$ wherein n is 1 to 4, Z is selected from the group consisting of F, Cl, Br, I, and mixtures thereof, and w ranges from 1 to 5, and aryl radicals

$$R^{16}$$
 R^{15}
 R^{19}

wherein R¹⁵, R¹⁶, R¹⁷, R¹⁸, and R¹⁹ are the same or different and arc selected from hydrogen and alkyl radicals having 1 to 20 carbon atoms, and mixtures thereof, to yield acyl sulfonate material of the general formula

$$R^{12} - C - O - SO_{2}R^{14}$$

$$R^{12} - C - O - SO_{2} - O - C - R^{14}L^{2}$$

$$R^{12} - C - O - SO_{2} - O - C - R^{14}L^{2}$$

$$R^{12} - C - O - SO_{2} - OR^{14}$$

$$R^{12} - C - O - SO_{2} - NR^{14}R^{14}$$

which is then reacted with a dioxane of the formula

$$\begin{array}{c}
R^{11} & O \\
R^{10} & O \\
R^{9} & O \\
R^{7}
\end{array}$$

wherein R⁴, R⁵, R⁶, R⁷, R⁸, R⁹, R¹⁰, and R¹¹ are the same or different and are selected from hydrogen, alkyl and hydroxyalkyl radicals having 1 to 4 carbons to yield

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which is then aminated with an alkylamine of the formula

$$H_2N - C - R^2$$

wherein R¹, R², and R³ are as previously defined to yield

$$R^{12} - \frac{\overset{O}{\overset{}{\parallel}}}{\overset{}{-}} \overset{R^{10}}{\overset{}{-}} \overset{R^8}{\overset{}{-}} \overset{R^8}{\overset{}{-}} \overset{R^6}{\overset{}{-}} \overset{R^4}{\overset{}{-}} \overset{R^1}{\overset{}{-}} \overset{R^1}{\overset{}{-}} \overset{R^2}{\overset{}{-}} \overset{R^{11}}{\overset{}{-}} \overset{R^9}{\overset{}{-}} \overset{R^8}{\overset{}{-}} \overset{R^8}{\overset{-$$

which is then hydrolyzed with base to yield

2. (original) The method of claim 1 for the synthesis of severely sterically hindered secondary aminoether alcohols using sulfonyl halide of the formula R¹⁴SO₂X.

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- 3. (original) The method of claim 1 for the synthesis of severely sterically hindered secondary aminoether alcohols using sulfuryl halide of the formula SO_2X_2 .
- 4. (original) The method of claim 1 for the synthesis of severely sterically hindered secondary aminoether alcohols using the mixed sulfuryl ester halide of the formula R¹⁴OSO₂X.
- 5. (original) The method of claim 1 for the synthesis of severely sterically hindered secondary aminoether alcohols using the mixed sulfuryl amide halide of the formula $R_2^{14,14}NSO_2X$.
- 6. (currently amended) The method of according to claim 1, 2, 3, 4 or 5 wherein R¹, R² and R³ are methyl radicals.
- 7. (currently amended) The method of according to claim 1, 2, 3, 4 or 5 wherein R⁴, R⁵, R⁶, R⁷, R⁸, R⁹, R¹⁰, and R¹¹ are hydrogen.
- 8. (currently amended) The method of according to claim 1, 2, 3, 4 or 5 wherein R¹⁵, R¹⁶, R¹⁸, and R¹⁹ are hydrogen and R¹⁷ is hydrogen or methyl.
- 9. (currently amended) The method of according to claim 1, 2, 3, 4 or 5 wherein the base is selected from alkali metal hydroxide, alkali metal alkoxide, or alkali metal carbonate.
- 10. (currently amended) The method of according to claim 1, 2, 3, 4 or 5 wherein Y is hydrogen or sodium.
- 11. (currently amended) The method of according to claim 1, 2, 3, 4 or 5 wherein R¹, R² and R³ are methyl, R⁴, R⁵, R⁶, R⁷, R⁸, R⁹, R¹⁰, and R¹¹ are hydrogen, R¹⁵, R¹⁶, R¹⁸, and R¹⁹ are hydrogen, R¹⁷ is hydrogen or methyl, and Y is hydrogen, sodium, or ammonium.

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- 12. (currently amended) The method of according to claim 1, 2, 3, 4 or 5 wherein the acyl sulfonate is made by reacting organic carboxylic acid or the salt of a carboxylic acid with the sulfonyl halide, sulfuryl halide, mixed sulfuryl ester halide or mixed sulfuryl amide halide at a temperature in the range of between about -20 to 200°C at a pressure between about 1 bar and 100 bars, the acyl sulfonate is reacted with the dioxane at a molar ratio of dioxane to acyl sulfonate in the range of 1:1 to 10:1 at a temperature of between about 50°C to about 200°C to yield a cleavage product, the cleavage product and the alkyl amine reacted at an amine to sulfonate group ratio ranging from about stoichiometric to about 10:1 at pressure of from about atmospheric (1 bar) to about 100 bars at temperature of from about 40°C to about 200°C, and the resulting aminated product is hydrolyzed with base at a temperature from about 20°C to about 110°C.
- 13. (currently amended) The method of according to claim 1, 2, 3, 4 or 5 wherein the organic carboxylic acid or the salt thereof, the sulfonyl halide, sulfuryl halide, mixed sulfuryl ester halide or mixed sulfuryl amide halide and the dioxane care combined in a single step to produce a reaction mixture, the reaction mixture being heated at a temperature of between about 50°C to about 200°C to produce the cleavage product, the cleavage product and the alkylamine are reacted at am amine to cleavage product ratio ranging from about stoichiometric to about 10:1 at a pressure from about atmospheric (1 bar) to about 100 bars at a temperature of from about 40°C to about 200°C, the resulting aminated product being reacted with base at a temperature from about 20°C to about 110°C.